Applicant : Nariyoshi Shinomiya et al.

For : c-met siRNA ADENOVIRUS VECTORS INHIBIT CANCER CELL

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## In the Claims:

This listing of claims will replace all prior versions and listings of claims in the application:

- (currently amended) An interfering RNA (RNAi) molecule having a sequence that is sufficiently complementary to the-a\_sequence of mRNA encoded by human c-met (SEQ ID NO:1), murine c-met (SEQ ID NO:2), or c-met of another mammalian source, so that expression of said RNAi molecule in a cell that normally expresses c-met results in diminution or loss of expression of said mRNA.
- (original) The RNAi molecule of claim 1 that is a single stranded siRNA that forms a hairpin structure.
- 3. (original) The RNAi molecule of claim 1 that is a double stranded siRNA.
- 4. (currently amended) The RNAi molecule of any of claims 1.3 claim 1 that (i) comprises, or (ii) hybridizes to a Met target sequence that comprises, a sequence selected from the group consisting of: (a) SEQ ID NO:9; (b) SEQ ID NO:10; (c) SEQ ID NO:11; (d) SEQ ID NO:12; (e) SEQ ID NO:13; (f) SEQ ID NO:14; (g) SEQ ID NO:15; (h) SEQ ID NO:16; (i) SEQ ID NO:17; and (j) SEQ ID NO:18.
- (currently amended) The RNAi molecule of any of claims 1.3 claim 1 that consists essentially of:
- (i) a sequence, selected from the group consisting of (a) SEQ ID NO:9; (b) SEQ ID NO:10; (c) SEQ ID NO:11; (d) SEQ ID NO:12; (e) SEQ ID NO:13; (f) SEQ ID NO:14; (g) SEQ ID NO:15; (h) SEQ ID NO:16; (i) SEQ ID NO:17; and (j) SEQ ID NO:18, or (ii) a sequence that hybrizes to a Met target selected from (a)- (j), above.

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 (original) The RNAi molecule of claim 4 that comprises a sequence complementary to human c-met mRNA which is selected from the group consisting of SEQ ID NO:13, SEQ ID NO:14, and SEQ ID NO:15.

- (original) The RNAi molecule of claim 5 that consists essentially of a sequence complementary to human c-met mRNA which is selected from the group consisting of SEQ ID NO:13, SEO ID NO:14, and SEO ID NO:15.
- (currently amended) An expression construct comprising DNA that encodes the RNAi
  molecule of any of claims 1.7claim 1 operatively linked to a promoter that drives the
  expression of said RNAi in a c-met-expressing cell.
- 10. (original) An expression construct comprising the DNA molecule of claim 8.
- (currently amended) The expression construct of claim 9-or-10, wherein the a promoter is one that drives the expression of said RNAi in a c-met-expressing tumor or cancer cell.
- 12. (currently amended) The expression construct of any of claims 9.11 claim 11 wherein the promoter is a pollII promoter.
- (original) The expression construct of claim 12 wherein the polIII promoter is a U6 promoter.
- 14. (currently amended) A viral vector comprising the expression construct of any of

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claims 9-13claim 9.

(currently amended) The viral vector of claim 14 that is a transient expression vector vector.

- 16. (original) The viral vector of claim 13 that is a stable expression vector.
- 17. (currently amended) The viral vector of claim 14 or 16-that is an adenoviral vector.
- 18. (original) The adenoviral vector of claim 17 that is an Ad5 viral vector.
- 19. (original) The Ad5 viral vector of claim 18 selected from the group consisting of: (a) si-mMet-Ad5<sup>50</sup>; (b) si-mMet-Ad5<sup>10</sup>; (c) si-mMet-Ad5<sup>10</sup>; (d) si-mMet-Ad5<sup>70</sup>; (e) si-hMet-Ad5<sup>16</sup>; (f) si-hMet-Ad5<sup>22</sup>; (h) si-dMet-Ad5<sup>111</sup>; (i) si-dMet-Ad5<sup>107</sup>; and (j) si-dMet-Ad5<sup>22</sup>.
- (original) The Ad5 viral vector of claim 19 wherein the vector is si-hMet-Ad5<sup>16</sup>; si-hMet-Ad5<sup>62</sup>; or si-hMet-Ad5<sup>221</sup>.
- 21-37. (canceled)
- 38. (currently amended) A method of treating a c-met\* tumor or cancer in a subject, comprising administering to the subject by an effective route, an amount of the viral vector of any-of-claims 14-20claim 14 effective for inhibiting expression of c-met and thereby (i) inhibiting the growth, invasion or metastasis of cells of said tumor or cancer, or (ii) killing said tumor or cancer cells.

39-47. (canceled)